IN THE CLAIMS

1. (Currently amended) A process for converting gaseous alkanes to liquid hydrocarbons comprising:

reacting a gaseous feed having lower molecular weight alkanes with bromine vapor to form alkyl bromides and hydrobromic acid; and

reacting said alkyl bromides in the presence of <u>said hydrobromic acid and a catalyst consisting essentially of</u> a synthetic crystalline alumino-silicate catalyst and at a temperature sufficient to form higher molecular weight hydrocarbons and additional hydrobromic acid.

- 2. (Currently amended) The process of claim 1 wherein said bromine <u>vapor</u> is substantially dry, thereby avoiding the formation of significant carbon dioxide along with said alkyl bromides.
- 3. (Original) The process of claim 1 wherein said gaseous feed is natural gas.
- 4. (Currently amended) The process of claim 3 wherein said natural gas is treated to remove substantially all of the carbon dioxide and sulfur compounds therefrom prior to reacting with said bromine <u>vapor</u>.
- 5. (Original) The process of claim 1 wherein said temperature is from about 150° C. to about 400° C.
- 6. (Original) The process of claim 5 wherein said temperature is from about 250° C. to about 350° C.
- 7. (Original) The process of claim 1 wherein said crystalline alumino-silicate catalyst is a zeolite catalyst.
- 8. (Original) The process of claim 7 wherein said zeolite catalyst is a ZSM-5 zeolite catalyst and said higher molecular weight hydrocarbons contain a C7+ fraction that is composed primarily of substituted aromatics.
- 9. (Original) The process of claim 8 wherein said ZSM-5 zeolite catalyst is modified with at least one modifying cation selected from hydrogen and Group IA alkaline metals, or Group IIA alkaline earth metals.

10. (Original) The process of claim 9 wherein said ZSM-5 zeolite catalyst is modified by ion exchange with at least one cation selected from hydrogen, sodium, potassium, cesium, magnesium, calcium or barium.

11. (Currently amended) The process of claim 22 18 further comprising wherein said water is an aqueous solution and said step of contacting comprises:

removing said hydrobromic acid and said additional hydrobromic acid from said higher molecular weight hydrocarbons by neutralization reaction with an said aqueous solution containing reaction products obtained by oxidizing an aqueous solution containing a metal bromide salt, the metal of said metal bromide salt being selected from Cu, Zn, Fe, Co, Ni, Mn, Ca or Mg bromide.

- 12. (Currently amended) The process of claim 1 wherein said bromine <u>vapor</u> is produced by oxidizing an aqueous metal bromide salt solution, the metal of said metal bromide salt being selected from Cu, Zn, Fe, Co, Ni, Mn, Ca, or Mg.
- 13. (Currently amended) The process of claim 22 18 further comprising wherein said step of contacting comprises:

removing said hydrobromic acid and said additional hydrobromic acid from said higher molecular weight hydrocarbons by dissolution into said <u>water</u> forming a hydrobromic acid solution, said hydrobromic acid solution being neutralized by reaction with an aqueous solution containing a metal hydroxide obtained by oxidizing an aqueous metal bromide salt solution with oxygen, the metal of said metal bromide salt being selected from Cu, Zn, Fe, Co, Ni, Mn, Ca or Mg.

14. (Currently amended) The process of claim 22 18 wherein said step of contacting comprises removing said hydrobromic acid and said additional hydrobromic acid are is removed from said higher molecular weight hydrocarbons by dissolution into said water forming a hydrobromic acid solution, said process further comprising:

<u>vaporizing said</u> hydrobromic acid solution being vaporized and reacted reacting said vaporized hydrobromic acid solution with a metal oxide, said metal oxide being obtained by oxidizing a metal bromide salt contained on a porous

support, the metal of said metal bromide salt being selected from the group Cu, Zn, Fe, Co, Ni, Mn, Ca or Mg.

- 15. (Currently amended) The process of claim 1 wherein said bromine <u>vapor</u> is produced by oxidizing a metal bromide salt contained on a porous support, the metal of said metal bromide salt being selected from the group Cu, Zn, Fe, Co, Ni, Mn, Ca, or Mg.
- 16. (Original) The process of Claim 1 wherein said higher molecular weight hydrocarbons contains a C3, C4 and C5+ fractions in admixture with excess lower alkanes, the process further comprising:

dehydrating said higher molecular weight hydrocarbons to a dew point of about -20° C. or less so as to recover said C5+ fractions as a liquid.

17. (Currently amended) The process of claim 16 further comprising:

mixing at least a portion of said C3 and C4 fractions with said alkyl bromides and said hydrobromic acid prior to the step of reacting over said synthetic crystalline alumino-silicate catalyst.

18. (Currently amended) A process for converting gaseous lower molecular weight alkanes to liquid hydrocarbons comprising:

reacting a gaseous feed containing lower molecular weight alkanes with bromine <u>vapor</u> to form alkyl bromides and hydrobromic acid;

reacting said alkyl bromides in the presence of a synthetic crystalline aluminosilicate catalyst <u>and hydrobromic acid</u> to form higher molecular weight hydrocarbons and additional hydrobromic acid; and

converting contacting said hydrobromic acid and, said additional hydrobromic acid to bromine, and said higher molecular weight hydrocarbons with water to remove said hydrobromic acid and additional hydrobromic acid from said higher molecular weight hydrocarbons.

- 19. (Currently amended) The process of claim 18 32 further comprising: dehydrating said higher molecular weight hydrocarbons.
- 20. (Currently amended) The process of claim 18 32 further comprising:

recycling said bromine that is converted from said hydrobromic acid and additional hydrobromic acid to said step of reacting with said gaseous feed, said bromine being recycled as a vapor.

- 21. (Currently amended) The process according to claim 1 wherein said step of reacting said gaseous feed with said bromine <u>vapor</u> occurs in a first reactor and said step of reacting said alkyl bromides occurs in a second reactor.
- 22. (Canceled)
- 23. (Canceled)
- 24. (Currently amended) The process of claim 1 wherein each of said bromine, said hydrobromic acid and said additional hydrobromic acid is a vapor.
- 25. (Currently amended) The process according to claim 18 wherein said step of reacting said gaseous feed with said bromine <u>vapor</u> occurs in a first reactor and said step of reacting said alkyl bromides occurs in a second reactor.
- 26. (Canceled)
- 27. (Canceled)
- 28. (Currently amended) The process of claim 18 wherein each of said bromine, said hydrobromic acid and said additional hydrobromic acid is a vapor.
- 29. (Currently amended) A process for converting gaseous alkanes to liquid hydrocarbons comprising:

Introducing introducing a mixture comprising lower molecular weight gaseous alkanes and bromine <u>vapor</u> to a first reactor;

withdrawing alkyl bromides and hydrobromic acid from said first reactor;

Introducing introducing said alkyl bromides and said hydrobromic acid into a second reactor containing a synthetic crystalline alumino-silicate catalyst; and

withdrawing <u>an effluent comprising</u> higher molecular weight hydrocarbons <u>and</u> <u>said hydrobromic acid from said second reactor.</u>

- 30. (Previously presented) The process of claim 29 wherein said higher molecular weight hydrocarbons contains C5+ fractions.
- 31. (Currently amended) The process of claim 29 wherein said bromine is a vapor and said hydrobromic acid is a vapor.
- 32. (New) The process of claim 18 further comprising:

converting said hydrobromic acid and said additional hydrobromic acid to bromine.